FDA 510(k) Premarket Notification

K896544 MD2 PERSONAL INFUSION SYSTEM

CLEARED

May 2, 1990
166 days to decision

K896544 · Product code: **FRN** · General Hospital
Source: https://www.510kdatabase.net/k896544/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pump, Infusion (FRN)
Date received	Nov 17, 1989
Decision date	May 2, 1990
Days to decision	166 days
Third-party review	No

APPLICANT

Company	**Norfolk Medical Products, Inc.**
Location	Walker, MI, US
Contact	MICHAEL J DALTON
510(k) history	20 submissions · 20 cleared · 1983-2020